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Tota	al Number of Pages in This Submission	59 / Montel South Manager 1700A1								
ENCLOSURES (Check all that apply)										
	Fee Transmittal Form Fee Attached Amendment/Reply After Final Affidavits/declaration(s) Extension of Time Request Express Abandonment Request Information Disclosure Statement Certified Copy of Priority Document(s) Response to Missing Parts/ Incomplete Application Response to Missing Parts under 37 CFR 1.52 or 1.53	Drawing(s) Licensing-related Papers Petition Petition to Convert to a Provisional Application Change of Correspondence Address Terminal Disclaimer Request for Refund CD, Number of CD(s) Appeal Communication to Board of Appeals and Interferences Appeal Communication to TC (Appeal Notice, Brief, Reply Brief) Proprietary Information Status Letter Other Enclosure(s) (please Identify below): Remarks Appeal Brief filed in triplicate								
		URE OF APPLICANT, ATTORNEY, OR AGENT								
Firm or Individ	Krisanne Shideler, Reg. N BLK Law Group	o. 36,272								
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FEE TRANSMITTAL for FY 2004

Effective 10/01/2003. Patent fees are subject to annual revision.

Applicant claims small entity status. See 37 CFR 1.27

TOTAL AMOUNT OF PAYMENT

(\$) 330.

Complete if Known							
Application Number	10/047,527						
Filing Date	10/23/2001	-					
First Named Inventor	Anthony M. Chasser						
Examiner Name	Melanie Bissett						
Art Unit	1711						
Attorney Docket No.	1700A1						

METHOD OF PAYMENT (check all that apply)	FEE CALCULATION (continued)					
Check Credit card Money Other None	3. ADDITIONAL FEES					
Deposit Account:	l -	Large Entity Small Entity				
Deposit 50000	Fee Code	Fee (\$)	Fee Code	Fee (\$)	Fee Description	Fee Paid_
Account Number 502800	1051	130	2051	65	Surcharge - late filing fee or oath	
Deposit Account BLK Law Group	1052	50	2052	25	Surcharge - late provisional filing fee or	
Name	1053	130	1053	130	cover sheet Non-English specification	
The Director is authorized to: (check all that apply) Charge fee(s) indicated below Credit any overpayments	1812	2,520	1812		For filing a request for ex parte reexamination	
Charge any additional fee(s) or any underpayment of fee(s)	1804	920*	1804	920*	Requesting publication of SIR prior to	
Charge fee(s) indicated below, except for the filing fee	1805	1,840*	1805	1 840*	Examiner action Requesting publication of SIR after	
to the above-identified deposit account.	1003	1,040	1000	1,040	Examiner action	
FEE CALCULATION	1251	110	2251	55	Extension for reply within first month	
1. BASIC FILING FEE	1252	420	2252	210	Extension for reply within second month	
Large Entity Small Entity Fee Fee Fee Fee Description Fee Paid	1253	950	2253		Extension for reply within third month	
Code (\$) Code (\$)		1,480	2254	740	Extension for reply within fourth month	
1001 770 2001 385 Utility filing fee	1255	2,010	2255	1,005	Extension for reply within fifth month	
1002 340 2002 170 Design filing fee	1401	330	2401		Notice of Appeal	
1003 530 2003 265 Plant filing fee	1402	330	2402		Filing a brief in support of an appeal	330
1004 770 2004 385 Reissue filing fee	1403	290	2403	145	Request for oral hearing	—
1005 160 2005 80 Provisional filing fee		1,510	1451		Petition to institute a public use proceeding	
SUBTOTAL (1) (\$)	1452	110	2452	55	Petition to revive - unavoidable	
2. EXTRA CLAIM FEES FOR UTILITY AND REISSUE		1,330	2453		Petition to revive - unintentional	
Fee from		1,330	2501		Utility issue fee (or reissue)	
Extra Claims below Fee Paid Total Claims -20** = X =	1502	480 640	2502		Design issue fee	
Independent - 3** = X = =	1503 1460	130	2503 1460		Plant issue fee Petitions to the Commissioner	
Multiple Dependent	1807	50	1807		Processing fee under 37 CFR 1.17(g)	
Large Entity Small Entity	1806	180	1806		Submission of Information Disclosure Stmt	
Fee Fee Fee <u>Fee Description</u> Code (\$) Code (\$)					Recording each patent assignment per	
1202 18 2202 9 Claims in excess of 20	8021	40	8021		property (times number of properties)	
1201 86 2201 43 Independent claims in excess of 3	1809	770	2809	385	Filing a submission after final rejection (37 CFR 1.129(a))	
1203 290 2203 145 Multiple dependent claim, if not paid	1810	770	2810	385	For each additional invention to be	
1204 86 2204 43 ** Reissue independent claims					examined (37 CFR 1.129(b))	
over original patent	1801	770	2801	385	()	
1205 18 2205 9 ** Reissue claims in excess of 20 and over original patent	1802	900	1802	900	Request for expedited examination of a design application	
SUBTOTAL (2) (\$)		Other fee (specify)				<u> </u>
**or number previously paid, if greater; For Reissues, see above	*Redu	iced by	Basic I	Filing F	ee Paid SUBTOTAL (3) (\$) 330.	

SUBMITTED BY (Complete (if applicable)) Registration No. Name (Print/Type) Krisanne Shideler 36,272 Telephone 724-934-5450 Signature 01/28/2004

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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of

: Confirmation No. 3962

Anthony M. Chasser et al.

: PATENT APPLICATION

Serial No. 10/047,527

: Group Art Unit: 1711

Filed: October 23, 2001

Examiner: Melanie Bissett

THE USE OF ANTI-OXIDANTS : Atty. Docket No. 1700A1

IN CLEAR POWDER COATINGS:

TO REDUCE FILIFORM

CORROSION OVER ALUMINUM:

BRIEF ON APPEAL

Mail Stop Appeal Brief - Patents Commissioner for Patents P. O. Box 1450 Alexandria, VA 22313-1450

Sir:

This is in support of the Notice of Appeal mailed November 26, 2003, and received in the U.S. Patent and Trademark Office on November 28, 2003, appealing the final rejection of claims 5-7, 9, 13-15, and 18. Commissioner is hereby authorized to charge deposit account 502800 for any and all fees necessary for filing this brief. The following headings correspond to the requirements of 37 CFR §1.192(c).

02/03/2004 WABDELR1 00000134 502800 10047527 01 FC:1401

330,00 DA

I I hereby certify that this correspondence is being deposited with the United States Postal Service as first class mail in an envelope addressed to: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450, on <u>January 28, 2004</u> (Date of Deposit and Signatur)

Krisanne Shid ler

(Signature)

(1) REAL PARTY IN INTEREST

The present application is assigned to PPG Industries Ohio, Inc. and the real party in interest is PPG Industries Ohio, Inc. having offices at 3800 West 143 Street, Cleveland, Ohio 44111, a wholly-owned subsidiary of PPG Industries, Inc. having offices at One PPG Place, Pittsburgh, Pennsylvania 15272.

(2) RELATED APPEALS AND INTERFERENCES

There are no related appeals or interferences known to appellants, the appellants' legal representative, or assignee which will directly affect or be directly affected by or have a bearing on the Board of Appeals ("Board")'s decision in the pending appeal.

(3) STATUS OF CLAIMS

Claims 5-7, 9, 13-15, and 18 are all the claims pending in the application, and all were rejected. Claims 1-4, 8, 10-12, 16, and 17 have been cancelled. Claims 3 (now cancelled), 5, 6, 9, 13-15, and 18 were rejected under 35 U.S.C. §102(b) as being unpatentable over Geary et al., U. S. Patent No. 4,801,680 (hereinafter referred to as "Geary") as evidenced by Laver, U. S. Patent No. 6,103,794 (hereinafter referred to as "Laver"). Claims 3, 5, 6, 13-15. and 18 were rejected under 35 U.S.C. §102(b) as being unpatentable over Chasser et al., U. S. Patent No. 6,069,221 (hereinafter referred to as "Chasser") as evidenced by Laver. Claim 4 (now cancelled), the subject matter of which has been incorporated into claim 18, was rejected under 35 U.S.C. §103(a) as being unpatentable over Chasser as evidenced by Laver, and in view of Nakae et al., U. S. Patent No. 5,719,212 (hereinafter referred to as "Nakae"). Claim 4, (again, the subject matter of which has been incorporated into claim 18), was additionally rejected under 35 U.S.C. §103(a) as being unpatentable over Geary as evidenced by Laver and in view of Nakae. Claim 7 was rejected under 35 U.S.C. §103(a) as being unpatentable over Geary as evidenced by Laver. Claims 5-7, 9, 13-15, and 18 are on appeal.

As discussed below, the amendment to claim 18 adding the language of cancelled claim 4 is believed to alter the basis of rejection of the remaining

claims. It is believed that the Examiner is now rejecting claims 5-7, 9, 13-15, and 18 under 35 U.S.C. §103(a) as being unpatentable over Geary as evidenced by Laver taken in view of Nakae. In a similar fashion, it is believed that the Examiner is now rejecting claims 5, 6, 13-15, and 18 under 35 U.S.C. §103(a) as being unpatentable over Chasser as evidenced by Laver taken in view of Nakae.

(4) STATUS OF AMENDMENTS

A Final Rejection in the above application was mailed August 26, 2003; claims 3-7, 9, 13-15, and 18, which includes all the claims currently pending in the application, were rejected. Claim 4 was rejected under 35 USC §112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which Applicant regards as the invention. In particular, claim 4 depended from a non-existent claim 19. Claim 4 was subsequently cancelled, the subject matter of which having been incorporated into independent claim 18, rendering the rejection under 35 USC §112, second paragraph, moot. Claims 3, 5, 6, 9, 13-15, and 18 were rejected under 35 U.S.C. §102(b) as being unpatentable over Geary as evidenced by Laver. Claims 3, 5, 6, 13-15, and 18 were rejected under 35 U.S.C. §102(b) as being unpatentable over Chasser as evidenced by Laver. Claim 4 was rejected under 35 U.S.C. §103(a) as being unpatentable over Chasser as evidenced by Laver, and in view of Nakae. As noted above it is assumed that this rejection will essentially be applied by the Examiner against claims 5, 6, 13-15, and 18. Claim 4 was additionally rejected under 35 U.S.C. §103(a) as being unpatentable over Geary as evidenced by Laver and in view of Nakae. As noted above it is assumed that this rejection will essentially be applied by the Examiner against claims 5-7, 9 13-15, and 18. Claim 7 was also rejected under 35 U.S.C. §103(a) as being unpatentable over Geary as evidenced by Laver, and this rejection was listed separately in the Office Action.

An Amendment After Final Action (hereinafter "Amendment") was timely filed. The Amendment cancelled claims 3 and 4 and substantively amended claim 18 by incorporating the subject matter of claim 4, to place the claims in

better condition for appeal. The Amendment was entered as evidenced in the Advisory Office Action dated December 9, 2003. The Examiner noted that the rejections were maintained, and it has been assumed by Appellants that they are applied as discussed above.

(5) SUMMARY OF THE INVENTION

The present invention provides a coated aluminum substrate containing a cured coating, which in turn is derived from a coating composition comprising: (a) a polymer containing carboxylic functional groups, (b) a betahydroxyalkylamide curing agent having functional groups reactive with the functional groups of the polymer which is present in an amount sufficient to cure the polymer, and (c) 0.5 to 10 weight percent (based on the total weight of resin solids in the coating composition) of 2,6-di-tert-butyl-4-methyl-phenol (page 2, lines 9-13; page 6, lines 26-31; page 8, line 9; claim 18). The coated substrate is explicitly characterized as having improved filiform corrosion resistance compared to a similar coated substrate that does not contain 2,6-ditert-butyl-4-methyl-phenol in the cured coating (page 11, Table 2, Example 1; claim 18). In one embodiment of the present invention, the polymer of (a) containing carboxylic functional groups may be selected from acrylic polymers, polyester polymers, and polyurethane polymers (page 2, lines 10-12; claim 5); in a particular embodiment, it is an acrylic polymer (claim 15). embodiment of the present invention, the polymer has a number average molecular weight of from 1000 to 20,000 (claim 6). The polymer additionally may have an equivalent weight of from 200 to 2500 (page 3, line 15; page 5, line 7; page 6, line 12; claim 7). The polymer of (a) may typically be present in the powder coating composition in an amount ranging from 10 to 80 percent by weight based on the total weight of resin solids in the powder coating composition (claim 13). In one embodiment of the invention, the betahydroxyalkylamide curing agent of (b) is bis-hydroxyethylamide (claim 9). The curing agent may be present in the powder coating composition in an amount ranging from 2 to 40 percent by weight based on the total weight of resin solids in the powder coating composition (page 7, lines 29-30; claim 14).

The coated aluminum substrates of the present invention, which contain a cured coating composition derived from a coating composition containing 2,6di-tert-butyl-4-methyl-phenol, demonstrate improved filiform corrosion resistance compared to coated aluminum substrates that are coated with compositions containing other phenolic anti-oxidants such as 3,5-di-tertiarybutyl-4-hydroxyphenol compounds. Filiform corrosion appears as a filamentous, worm-like defect under a coating layer, adversely affecting appearance, and worse, often leads to coating delamination (peeling), making it a serious problem. It is particularly common over aluminum. Table 2 on page 11 in the specification shows a comparison of a coated aluminum substrate of the present invention compared to substrates coated with coating compositions containing hydroxyl-adjacent di-substituted phenolic anti-oxidants (such as Irganox® 1076, a 3,5-di-tertiary-butyl-4-hydroxyphenol compound, also disclosed in Chasser and Geary). Note that the coated aluminum substrate prepared in accordance with the present invention; i. e., coated with compositions containing 2,6-di-tert-butyl-4-methyl-phenol, (Example 1, row 1 of Table 2) demonstrates markedly improved filiform corrosion resistance as evidenced by lower filament density compared to substrates coated with compositions containing other hydroxyl-adjacent di-substituted phenolic antioxidants (see Examples 1, 4, 5, and 6, where the substrates coated with compositions containing hydroxyl-adjacent di-substituted phenolic anti-oxidants are compared directly to the coated substrate of the present invention). The presence of 2,6-di-tert-butyl-4-methyl-phenol in the coating compositions used to prepare the coated aluminum substrates of the present invention is critical to the filiform corrosion resistance performance of the invention. The examples further demonstrate the use of 2,6-di-tert-butyl-4-methyl-phenol within the claimed ranges of 0.5 to 10 percent by weight based on the total weight of resin solids in the powder coating composition.

(6) <u>ISSUES</u>

(i) No issues relating to 35 U.S.C. §112, first paragraph.

(ii) No issues relating to 35 U.S.C. §112, second paragraph. The rejection of claim 4 under 35 U.S.C. §112, second paragraph, is now moot with the cancellation of claim 4.

(iii) No issues relating to 35 U.S.C. §102. The rejections under 35 U.S.C. §102 are believed to be moot with the incorporation of the subject matter of now cancelled claim 4 into independent claim 18.

(iv) There are two issues relating to 35 U.S.C. §103(a).

(a) Whether the subject matter of 5-7, 9, 13-15, and 18 are obvious in view of the teachings of Geary as evidenced by Laver taken in view of the teachings of Nakae.

(b) Whether the subject matter of 5, 6, 13-15, and 18 are obvious in view of the teachings of Chasser as evidenced by Laver taken in view of the teachings of Nakae.

(7) GROUPING OF CLAIMS

Claims 5, 6, 13, 14 and 18 stand or fall together. Claims 7 and 9 stand or fall together. Claims 7 and 9 do not have Chasser applied against them (Issue (iv) (b)) and therefore stand separately from 5, 6, 13, 14 and 18. Claim 15 stands or falls alone. Claim 15 includes limitations that further distinguish this claim from the teachings of Geary, which supports the standing of claim 15.

(8) **ARGUMENTS**

(i) No issue.

<u>(ii) No issue.</u>

(iii) No issue.

(iv) (a) Wh ther the subject matter f 5-7, 9, 13-15, and 18 are obvious in view of the tachings of Geary as videnced by Laver taken in view of the teachings of Nakae.

Claims 5, 6, 13, 14 and 18:

Claims 5, 6, 13, 14, and 18 are assumed to be rejected under 35 U.S.C. §103(a) as being obvious over Geary as evidenced by Laver and in view of Nakae. In both the Office Action of April 3, 2003, and the Final Office Action of August 26, 2003, the Examiner asserts that "Geary discloses thermosetting powder coating compositions comprising carboxylic acid-containing polyesters and beta-hydroxyalkylamide curing agents" (Geary, col. 1, lines 34-39). The Examiner further asserts that "examples show bis-hydroxyethylamide compounds and carboxylic acid group-containing polyester polymers having a number average molecular weight of 3500-4500. Example 1A suggests the combination of 72.4 percent by weight carboxylic acid-containing polyester, 4.3 percent by weight bis-hydroxyethylamide, and 1.5 percent by weight Irganox® 1076." Laver provides the structure of Irganox® 1076, which is n-octadecyl-3-(3,5-di-t-butyl-4-hydroxyphenyl)propionate, a phenolic compound having branched butyl groups in positions ortho to the hydroxy group. The coatings are applied to metal substrates, including aluminum. As noted by the Examiner in the Office Action of April 3, 2003, neither Geary nor Laver mentions the use of 2,6-di-tert-butyl-4-methyl-phenol. The Nakae reference teaches acid-epoxy type powder coating compositions comprising antioxidants having a melting point of from 50°C to 140°C. The Examiner relies on Nakae for teaching the use of 2,6-di-tert-butyl-4-methyl-phenol in powder coating compositions, asserting that certain phenol antioxidants, including 2,6-di-tert-butyl-4-methylphenol, are preferred in powder compositions because of their melting points. The Examiner concludes that it would have been obvious to one skilled in the art to choose 2,6-di-tert-butyl-4-methyl-phenol as a phenolic antioxidant in Geary's composition to form coatings having improved blocking resistance and melt processability.

Appellants respectfully submit that the basis on which the claims were rejected is not a valid rejection under 35 U. S. C. §103(a). First, if the references are considered each in their entirety, there is no motivation to combine the teachings of Geary with those of Nakae. Secondly, even if one were to combine the whole teaching of Geary with the whole teaching of Nakae, one would not arrive at the present invention as claimed.

References Considered as a Whole Offer No Motivation to Combine

The motivation to modify the prior art must flow from some teaching in the art that suggests the desirability or incentive to make the modification needed to arrive at the claimed invention. See In re Napier, 55 F.3d 610, 613, 34 U.S.P.Q.2d 1782, 1784 (Fed. Cir. 1995). Geary discloses thermosetting powder coating compositions that comprise a co-reactable particulate mixture of carboxylic acid group-containing polyesters and beta-hydroxyalkylamide curing agents (Abstract). Examples 1A, 2A, 3 and 4 disclose the use of Irganox® 1076 as an additive in these powder coating compositions. The additive is defined in footnote 3 of Example 1 as a polyphenol stabilizer that inhibits oxidation. Anti-oxidants are not taught, suggested, or recommended anywhere else in the reference. All of the examples in Geary demonstrate coatings applied over steel panels. Nakae discloses a different powder coating composition relying on a different cure chemistry, comprising an epoxy groupcontaining acrylic resin; a polycarboxylic acid; and an anti-oxidant having a melting point of from 50° to 140°C (Abstract). A list of over fifty anti-oxidants is provided (Nakae col. 3. Line 35-col. 4, line 49), which includes 2,6-di-tert-butyl-4-methyl-phenol and n-octadecyl-3-(3,5-di-t-butyl-4-hydroxyphenyl)propionate (Irganox® 1076). There is no teaching or suggestion in Nakae that the antioxidants used therein are suitable for use in any other coatings; moreover, coatings comprising carboxylic acid group-containing resins and betahydroxyalkylamide curing agents are not disclosed at all. The only coating composition disclosed in Nakae undergoes an acid-epoxy type cure mechanism. There is no teaching or suggestion that any of the anti-oxidants in Nakae would be effective in a carboxylic acid/beta-hydroxyalkylamide system such as that disclosed in Geary, much less that 2,6-di-tert-butyl-4-methyl-phenol would be equally effective as Irganox® 1076 in the system disclosed by Geary, such that one skilled in the art would be led to substitute the one anti-oxidant for the other.

The Examiner in both Office Actions alleges that the motivation to modify the composition of Geary with the 2,6-di-tert-butyl-4-methyl-phenol of Nakae is in order to form coatings having improved blocking resistance and melt processability. However, this proposed motivation to modify the Geary reference with the teaching of Nakae to use 2,6-di-tert-butyl-4-methyl-phenol is not supported. There is *no* teaching or suggestion in Nakae that 2,6-di-tert-butyl-4-methyl-phenol may be better than Irganox® 1076 for blocking resistance and melt processability in a coating composition that contains carboxylic acid functional polymers and beta-hydroxyalkylamide curing agents, such as that disclosed in Geary, or even that 2,6-di-tert-butyl-4-methyl-phenol is better than Irganox® 1076 for blocking resistance and melt processability in *any* powder coating composition.

There is absolutely no motivation to modify the compositions of Geary by particularly choosing the 2,6-di-tert-butyl-4-methyl-phenol of Nakae for improving blocking resistance and/or melt processability as suggested by the Examiner, or for any other reason, because none is offered by either reference, taken alone or in combination. Additionally, there is no teaching or suggestion in either reference, taken alone or in combination, to use the 2,6-di-tert-butyl-4-methyl-phenol of Nakae in a coating composition that contains carboxylic acid functional polymers and beta-hydroxyalkylamide curing agents, much less over an aluminum substrate, since neither reference demonstrates the use of such a combination over aluminum.

References Considered as a Whole Teach Away from Invention

If one were to combine the whole teachings of Geary with the whole teaching of Nakae, one would not arrive at the present invention as claimed; i. e., a coated aluminum substrate containing a cured coating derived from a coating composition comprising:

- (a) a polymer containing carboxylic functional groups;
- (b) a beta hydroxyalkylamide curing agent having functional groups reactive with the functional groups of the polymer which is present in an amount sufficient to cure the polymer; and
- (c) 0.5 to 10 weight percent based on the total weight of resin solids in the coating composition of 2,6-di-tert-butyl-4-methyl-phenol, the coated substrate being characterized as having improved filiform corrosion resistance compared to a similar coated substrate that does not contain (c) in the cured coating.

"It is impermissible within the framework of section 103 to pick and choose from any one reference only so much of it as will support a given position, to the exclusion of other parts necessary to the full appreciation of what such reference fairly suggests to one of ordinary skill in the art." In re Wesslau, 353 F.2d 238, 241, 147 U.S.P.Q. 391, 393 (C.C.P.A. 1965). See also MPEP §2141.02: "[a] prior art reference must be considered in its entirety, i. e., as a whole, including portions that would lead away from the claimed invention." If the references are considered each in their entirety, the teachings of Nakae combined with Geary teach away from the present invention. As mentioned earlier, Geary discloses thermosetting powder coating compositions that comprise a co-reactable particulate mixture of carboxylic acid groupcontaining polyesters and beta-hydroxyalkylamide curing agents. Nakae discloses a powder coating composition comprising an epoxy group-containing acrylic resin; a polycarboxylic acid; and any of over fifty anti-oxidants having a melting point of from 50° to 140°C. While a variety of metal substrates including both steel and aluminum are disclosed in Nakae (col. 6, lines 26-30), the only substrate on which the coating composition is demonstrated is steel. Likewise, all of the examples in Geary demonstrate coatings applied over steel panels. If one were to combine both references after considering them in their entirety, in order to prepare a coated substrate containing a cured coating, which in turn is derived from a coating composition comprising a polymer containing carboxylic functional groups, a beta-hydroxyalkylamide curing agent, and an anti-oxidant, one skilled in the art would use a steel substrate, since both references clearly demonstrate the use of their respective coating compositions over steel. Additionally, one is not likely to choose 2,6-di-tert-butyl-4-methyl-phenol from among a list of over fifty other anti-oxidants, without any guidance to do so offered by the Nakae reference. Further, since neither reference recognizes the improvement in filiform corrosion resistance over aluminum afforded by the use of 2,6-di-tert-butyl-4-methyl-phenol, there would be no particular impetus to use an aluminum substrate or 2,6-di-tert-butyl-4-methyl-phenol, and therefore it would not have been obvious to do so.

Claim 15

The discussion under the heading "Claims 5, 6, 13, 14, and 18" is incorporated herein by reference.

Claim 15 recites the coated aluminum substrate of claim 18 where said polymer is an acrylic polymer. The Examiner asserts in both Office Actions that Example 2A in Geary "shows the combination of carboxylic acid-containing polyester, carboxylic acid-containing acrylic polymer, a bis-hydroxyethylamide, and Irganox® 1076". As noted above, Geary discloses thermosetting powder coating compositions that comprise a co-reactable particulate mixture of carboxylic acid group-containing *polyesters* and beta-hydroxyalkylamide curing agents. The acrylic polymer disclosed in Example 2 is used in very minor amounts (less than 5 percent by weight of the resinous binder; less than 3.5 percent by weight of total resin solids). One skilled in the art would hardly look to Geary for a teaching of coating compositions containing carboxylic acid functional acrylics because Geary teaches a powder coating composition containing polyester polymers. Claim 15 is therefore believed to further distinguish the coated aluminum substrate of the present invention from Geary taken alone or in combination with Nakae.

Claims 7 and 9

The discussion under the heading "Claims 5, 6, 13, 14, and 18" is incorporated herein by reference.

Claim 7 recites the coated aluminum substrate of claim 18 wherein said polymer has an equivalent weight of from 200 to 2500. Claim 9 recites the

coated aluminum substrate of claim 18 wherein the beta-hydroxyalkylamide is bis-hydroxyethylamide. The arguments presented above with respect to claims 5, 6, 13, 14, and 18 apply here, and it is noted as above in the Grouping of Claims that claims 7 and 9 stand or fall together, independently of 5, 6, 13, 14 and 18. Claims 7 and 9 do not have Chasser applied against them (Issue (iv)(b)) and therefore stand separately from 5, 6, 13, 14 and 18.

(iv)(b) Whether the subject matter of 5, 6, 13-15, and 18 are obvious in view of the teachings of Chasser as evidenced by Laver taken in view of the teachings of Nakae.

Claims 5, 6, 13, 14 and 18:

Claims 5, 6, 13, 14, and 18 are assumed to be rejected under 35 U.S.C. §103(a) as being obvious over Chasser as evidenced by Laver and in view of Nakae. In both the Office Action of April 3, 2003, and the Final Office Action of August 26, 2003, the Examiner asserts that "Chasser discloses curable powder coating compositions comprising polymers with functional groups and curing agents reactive with the polymers that are applied to aluminum substrates (Abstract)." The Examiner further asserts that the polymers are selected from carboxylic acid group-containing polyester polymers having a number average molecular weight of 2000-3000, carboxylic acid group-containing acrylic polymers, carboxylic acid group-containing polyurethane polymers, and epoxy functional polymers. Polymers are used in preferred amounts of 50-85 percent by weight. Curing agents include beta-hydroxyalkylamides triglycidylisocyanurate (TGIC). Irganox® anti-oxidants are noted as suitable additives. Example A shows the combination of a carboxylic acid-containing polyester, 12.9 percent by weight TGIC, and about 1 percent by weight Irganox® 1076. Laver provides the structure of Irganox® 1076. Example C shows the combination of a carboxylic acid-containing polyester, a betahydroxyalkylamide curing agent, and Tinuvin® 900, a phenolic ultraviolet light stabilizer having substituents at ortho positions to the hydroxy group. As noted by the Examiner in the Office Action of April 3, 2003, neither Chasser nor Laver mentions the use of 2,6-di-tert-butyl-4-methyl-phenol. The Nakae reference teaches acid-epoxy type powder coating compositions comprising antioxidants having a melting point of from 50°C to 140°C. The Examiner relies on Nakae for teaching the use of 2,6-di-tert-butyl-4-methyl-phenol in powder coating compositions, asserting that certain phenol antioxidants, including 2,6-di-tert-butyl-4-methyl-phenol, are preferred in powder compositions because of their melting points. The Examiner concludes that it would have been obvious to one skilled in the art to choose 2,6-di-tert-butyl-4-methyl-phenol as a phenolic antioxidant in Chasser's composition to form coatings having improved blocking resistance and melt processability.

Appellants respectfully submit that the basis on which the claims were rejected is not a valid rejection under 35 U. S. C. §103(a). First, if the references are considered each in their entirety, there is no motivation to combine the teachings of Chasser with those of Nakae. Secondly, even if one were to combine the whole teaching of Chasser with the whole teaching of Nakae, one would not arrive at the present invention as claimed.

References Considered as a Whole Offer No Motivation to Combine

The Examiner in both Office Actions alleges that the motivation to modify the composition of Chasser with the 2,6-di-tert-butyl-4-methyl-phenol of Nakae is in order to form coatings having improved blocking resistance and melt processability. However, this proposed motivation to modify the Chasser reference with the teaching of Nakae to use 2,6-di-tert-butyl-4-methyl-phenol is not supported. As noted above, there is *no* teaching or suggestion in Nakae that 2,6-di-tert-butyl-4-methyl-phenol is better than Irganox® 1076 for blocking resistance and melt processability in *any* powder coating composition, let alone that 2,6-di-tert-butyl-4-methyl-phenol may be better than Irganox® 1076 for blocking resistance and melt processability in a coating composition that contains carboxylic acid functional polymers and beta-hydroxyalkylamide curing agents, which is only one of a collection of coating systems disclosed by Chasser that includes carboxylic acid functional polymers/polyepoxide curing

agents, carboxylic acid functional polymers/TGIC, and epoxy functional polymers/polyacid curing agents.

One skilled in the art would not be motivated to improve blocking resistance and/or melt processability of the Chasser composition by particularly choosing the 2,6-di-tert-butyl-4-methyl-phenol of Nakae because no teaching or suggestion is made in either reference, taken alone or in combination, of improved blocking resistance and/or melt processability offered by 2,6-di-tert-butyl-4-methyl-phenol. Additionally, there is no teaching or suggestion in either reference, taken alone or in combination, to select the 2,6-di-tert-butyl-4-methyl-phenol of Nakae for any reason and use it in a coating composition that contains carboxylic acid functional polymers and beta-hydroxyalkylamide curing agents.

References Considered as a Whole Teach Away from Invention

If one were to combine the whole teachings of Chasser with the whole teaching of Nakae, one would not arrive at the present invention as claimed. Chasser discloses curable powder coating compositions comprising polymers with functional groups and curing agents reactive with the polymers. The composition further comprises an oligomeric acid functional polyester with a particular structure as shown in col. 2, lines 40-48, which is the defining feature of the Chasser composition. The oligomeric polyester provides the composition with improved adhesion to metal and filiform corrosion resistance. The polymers with functional groups used by Chasser may be any of a variety of polymers with any of a variety of functional groups. Likewise, the curing agents may be selected from any of a number of different materials, depending on the functionality of the polymer. As mentioned above, carboxylic acid functional polymers/polyepoxide curing agents, carboxylic acid functional polymers/TGIC, and epoxy functional polymers/polyacid curing agents are all disclosed by Chasser as suitable coating systems. Nakae discloses exclusively an epoxy group-containing acrylic resin; a polycarboxylic acid; and an anti-oxidant having a melting point of from 50° to 140°C (Abstract). As noted above, list of over fifty anti-oxidants is provided, which includes 2,6-ditert-butyl-4-methyl-phenol and Irganox® 1076. There is no teaching or suggestion in Nakae that the anti-oxidants used therein are suitable for use in any other coatings; moreover, coatings comprising carboxylic acid group-containing resins and beta-hydroxyalkylamide curing agents are not disclosed at all in Nakae. The only coating composition disclosed in Nakae undergoes an acid-epoxy type cure mechanism. There is no teaching or suggestion that any of the other anti-oxidants in Nakae would be effective in a carboxylic acid/beta-hydroxyalkylamide system, much less that 2,6-di-tert-butyl-4-methyl-phenol would be equally effective as Irganox® 1076 a carboxylic acid/beta-hydroxyalkylamide system, such that one skilled in the art would be led to use the particular anti-oxidant 2,6-di-tert-butyl-4-methyl-phenol in the particular coating composition comprising carboxylic acid group-containing resins and beta-hydroxyalkylamide curing agents.

If the references are considered each in their entirety, the teachings of Nakae combined with Chasser teach away from the present invention. As mentioned earlier, Chasser discloses curable powder coating compositions comprising polymers with any of a number of functional groups and curing agents reactive with the polymers. Nakae discloses a powder coating composition comprising an epoxy group-containing acrylic resin; a polycarboxylic acid; and any of over fifty anti-oxidants having a melting point of from 50° to 140°C. If one were to combine both references after considering them in their entirety, in order to prepare a coated substrate containing a cured coating, which in turn is derived from a coating composition comprising a polymer containing functional groups, a curing agent, and an anti-oxidant, one skilled in the art would use an epoxy group-containing resin and a polycarboxylic acid curing agent, since that is the coating composition common to both references. Additionally, one is not likely to choose 2,6-di-tert-butyl-4methyl-phenol from among a list of over fifty other anti-oxidants, without any guidance to do so offered by the Nakae reference. Further, since neither reference recognizes the improvement in filiform corrosion resistance over aluminum afforded by the use of 2,6-di-tert-butyl-4-methyl-phenol (Chasser has already achieved filiform corrosion resistance with the oligomeric polyester),

there would be no particular reason to use 2,6-di-tert-butyl-4-methyl-phenol, and therefore it would not have been obvious to do so.

Claim 15

The discussion under the heading "Claims 5, 6, 13, 14, and 18" is incorporated herein by reference.

Claim 15 recites the coated aluminum substrate of claim 18 where said polymer is an acrylic polymer. The arguments presented above with respect to claims 5, 6, 13, 14, and 18 apply here, and it is noted as above in the Grouping of Claims that claim 15 includes limitations that further distinguish this claim from the teachings of Geary (Issue (iv)(a)), which supports the standing of claim 15.



CONCLUSION

Appellants have shown that if the references are considered each in their entirety, there is no motivation to combine the teachings of the Geary reference with Nakae in order to arrive at the present invention. Likewise, there is no motivation to combine the teachings of the Chasser reference with Nakae in order to arrive at the present invention. Appellants have further shown that if one were to combine the whole teachings of Geary or Chasser with the whole teaching of the Nakae, one would not arrive at the present invention as claimed.

Based on the above, Appellants respectfully request that the Board reverse the Examiner on the rejections of claims 5, 6, 7, 9, 13, 14, 15 and 18 as unpatentable under 35 U.S.C. 103 over the applied prior art and allow these claims.

Respectfully submitted,

KRISANNE SHIDELER Registration No. 36,272 Agent of Record

(724) 934-5450

Pittsburgh, Pennsylvania January 28, 2004

App ndix THE CLAIMS ON APPEAL

- 5. The coated aluminum substrate of claim 18 wherein said polymer containing carboxylic groups is selected from the group consisting of acrylic polymers, polyester polymers, and polyurethane polymers.
- 6. The coated aluminum substrate of claim 18 wherein said polymer has a number average molecular weight of from 1,000 to 20,000.
- 7. The coated aluminum substrate of claim 18 wherein said polymer has an equivalent weight equal from 200 to 2,500.
- 9. The coated aluminum substrate of claim 18 wherein the betahydroxyalkylamide is bis-hydroxyethylamide.
- 13. The coated aluminum substrate of claim 18 wherein said polymer is present in an amount ranging from 10 to 80 weight percent based on the total weight resin solids in the powder coating composition.
- 14. The coated aluminum substrate of claim 18 wherein said curing agent is present in an amount ranging from 2 to 40 weight percent based on the total weight resin solids in the powder coating composition.

- 15. The coated aluminum substrate of claim 18 where said polymer is an acrylic polymer.
- 18. A coated aluminum substrate containing a cured coating derived from a coating composition comprising:
 - a polymer containing carboxylic functional groups;
- b. a beta hydroxyalkylamide curing agent having functional groups reactive with the functional groups of the polymer which is present in an amount sufficient to cure the polymer; and
- c. 0.5 to 10 weight percent based on the total weight of resin solids in the coating composition of 2,6-di-tert-butyl-4-methyl-phenol, the coated substrate being characterized as having improved filiform corrosion resistance compared to a similar coated substrate that does not contain (c) in the cured coating.



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of

: Confirmation No. 3962

Anthony M. Chasser et al.

: PATENT APPLICATION

Serial No. 10/047,527

: Group Art Unit: 1711

Filed: October 23, 2001

: Examiner: Melanie Bissett

THE USE OF ANTI-OXIDANTS : Atty. Docket No. 1700A1

IN CLEAR POWDER COATINGS: TO REDUCE FILIFORM

CORROSION OVER ALUMINUM:

BRIEF ON APPEAL

Mail Stop Appeal Brief - Patents Commissioner for Patents P. O. Box 1450 Alexandria, VA 22313-1450

Sir:

This is in support of the Notice of Appeal mailed November 26, 2003, and received in the U.S. Patent and Trademark Office on November 28, 2003, appealing the final rejection of claims 5-7, 9, 13-15, and 18. Commissioner is hereby authorized to charge deposit account 502800 for any and all fees necessary for filing this brief. The following headings correspond to the requirements of 37 CFR §1.192(c).

> I hereby certify that this correspondence is being deposited with the United States Postal Service as first class mail in an envelope addressed to: Commissioner for Patents, P.O. Box 1450, Alexandria, VA 22313-1450, on <u>January 28, 2004</u> (Date of Deposit and Signature) Krisann Shideler (Signature)

(1) REAL PARTY IN INTEREST

The present application is assigned to PPG Industries Ohio, Inc. and the real party in interest is PPG Industries Ohio, Inc. having offices at 3800 West 143 Street, Cleveland, Ohio 44111, a wholly-owned subsidiary of PPG Industries, Inc. having offices at One PPG Place, Pittsburgh, Pennsylvania 15272.

(2) RELATED APPEALS AND INTERFERENCES

There are no related appeals or interferences known to appellants, the appellants' legal representative, or assignee which will directly affect or be directly affected by or have a bearing on the Board of Appeals ("Board")'s decision in the pending appeal.

(3) STATUS OF CLAIMS

Claims 5-7, 9, 13-15, and 18 are all the claims pending in the application, and all were rejected. Claims 1-4, 8, 10-12, 16, and 17 have been cancelled. Claims 3 (now cancelled), 5, 6, 9, 13-15, and 18 were rejected under 35 U.S.C. §102(b) as being unpatentable over Geary et al., U. S. Patent No. 4,801,680 (hereinafter referred to as "Geary") as evidenced by Laver, U. S. Patent No. 6,103,794 (hereinafter referred to as "Laver"). Claims 3, 5, 6, 13-15, and 18 were rejected under 35 U.S.C. §102(b) as being unpatentable over Chasser et al., U. S. Patent No. 6,069,221 (hereinafter referred to as "Chasser") as evidenced by Laver. Claim 4 (now cancelled), the subject matter of which has been incorporated into claim 18, was rejected under 35 U.S.C. §103(a) as being unpatentable over Chasser as evidenced by Laver, and in view of Nakae et al., U. S. Patent No. 5,719,212 (hereinafter referred to as "Nakae"). Claim 4, (again, the subject matter of which has been incorporated into claim 18), was additionally rejected under 35 U.S.C. §103(a) as being unpatentable over Geary as evidenced by Laver and in view of Nakae. Claim 7 was rejected under 35 U.S.C. §103(a) as being unpatentable over Geary as evidenced by Laver. Claims 5-7, 9, 13-15, and 18 are on appeal.

As discussed below, the amendment to claim 18 adding the language of cancelled claim 4 is believed to alter the basis of rejection of the remaining

claims. It is believed that the Examiner is now rejecting claims 5-7, 9, 13-15, and 18 under 35 U.S.C. §103(a) as being unpatentable over Geary as evidenced by Laver taken in view of Nakae. In a similar fashion, it is believed that the Examiner is now rejecting claims 5, 6, 13-15, and 18 under 35 U.S.C. §103(a) as being unpatentable over Chasser as evidenced by Laver taken in view of Nakae.

(4) STATUS OF AMENDMENTS

A Final Rejection in the above application was mailed August 26, 2003; claims 3-7, 9, 13-15, and 18, which includes all the claims currently pending in the application, were rejected. Claim 4 was rejected under 35 USC §112, second paragraph, as being indefinite for failing to particularly point out and distinctly claim the subject matter which Applicant regards as the invention. In particular, claim 4 depended from a non-existent claim 19. Claim 4 was subsequently cancelled, the subject matter of which having been incorporated into independent claim 18, rendering the rejection under 35 USC §112, second paragraph, moot. Claims 3, 5, 6, 9, 13-15, and 18 were rejected under 35 U.S.C. §102(b) as being unpatentable over Geary as evidenced by Laver. Claims 3, 5, 6, 13-15, and 18 were rejected under 35 U.S.C. §102(b) as being unpatentable over Chasser as evidenced by Laver. Claim 4 was rejected under 35 U.S.C. §103(a) as being unpatentable over Chasser as evidenced by Laver, and in view of Nakae. As noted above it is assumed that this rejection will essentially be applied by the Examiner against claims 5, 6, 13-15, and 18. Claim 4 was additionally rejected under 35 U.S.C. §103(a) as being unpatentable over Geary as evidenced by Laver and in view of Nakae. As noted above it is assumed that this rejection will essentially be applied by the Examiner against claims 5-7, 9 13-15, and 18. Claim 7 was also rejected under 35 U.S.C. §103(a) as being unpatentable over Geary as evidenced by Laver, and this rejection was listed separately in the Office Action.

An Amendment After Final Action (hereinafter "Amendment") was timely filed. The Amendment cancelled claims 3 and 4 and substantively amended claim 18 by incorporating the subject matter of claim 4, to place the claims in

better condition for appeal. The Amendment was entered as evidenced in the Advisory Office Action dated December 9, 2003. The Examiner noted that the rejections were maintained, and it has been assumed by Appellants that they are applied as discussed above.

(5) SUMMARY OF THE INVENTION

The present invention provides a coated aluminum substrate containing a cured coating, which in turn is derived from a coating composition comprising: (a) a polymer containing carboxylic functional groups, (b) a betahydroxyalkylamide curing agent having functional groups reactive with the functional groups of the polymer which is present in an amount sufficient to cure the polymer, and (c) 0.5 to 10 weight percent (based on the total weight of resin solids in the coating composition) of 2,6-di-tert-butyl-4-methyl-phenol (page 2, lines 9-13; page 6, lines 26-31; page 8, line 9; claim 18). The coated substrate is explicitly characterized as having improved filiform corrosion resistance compared to a similar coated substrate that does not contain 2,6-ditert-butyl-4-methyl-phenol in the cured coating (page 11, Table 2, Example 1; claim 18). In one embodiment of the present invention, the polymer of (a) containing carboxylic functional groups may be selected from acrylic polymers, polyester polymers, and polyurethane polymers (page 2, lines 10-12; claim 5); in a particular embodiment, it is an acrylic polymer (claim 15). In one embodiment of the present invention, the polymer has a number average molecular weight of from 1000 to 20,000 (claim 6). The polymer additionally may have an equivalent weight of from 200 to 2500 (page 3, line 15; page 5, line 7; page 6, line 12; claim 7). The polymer of (a) may typically be present in the powder coating composition in an amount ranging from 10 to 80 percent by weight based on the total weight of resin solids in the powder coating composition (claim 13). In one embodiment of the invention, the betahydroxyalkylamide curing agent of (b) is bis-hydroxyethylamide (claim 9). The curing agent may be present in the powder coating composition in an amount ranging from 2 to 40 percent by weight based on the total weight of resin solids in the powder coating composition (page 7, lines 29-30; claim 14).

The coated aluminum substrates of the present invention, which contain a cured coating composition derived from a coating composition containing 2,6filiform corrosion demonstrate improved di-tert-butyl-4-methyl-phenol, resistance compared to coated aluminum substrates that are coated with compositions containing other phenolic anti-oxidants such as 3,5-di-tertiary-Filiform corrosion appears as a butyl-4-hydroxyphenol compounds. filamentous, worm-like defect under a coating layer, adversely affecting appearance, and worse, often leads to coating delamination (peeling), making it a serious problem. It is particularly common over aluminum. Table 2 on page 11 in the specification shows a comparison of a coated aluminum substrate of the present invention compared to substrates coated with coating compositions containing hydroxyl-adjacent di-substituted phenolic anti-oxidants (such as Irganox® 1076, a 3,5-di-tertiary-butyl-4-hydroxyphenol compound, also disclosed in Chasser and Geary). Note that the coated aluminum substrate prepared in accordance with the present invention; i. e., coated with compositions containing 2,6-di-tert-butyl-4-methyl-phenol, (Example 1, row 1 of Table 2) demonstrates markedly improved filiform corrosion resistance as evidenced by lower filament density compared to substrates coated with compositions containing other hydroxyl-adjacent di-substituted phenolic antioxidants (see Examples 1, 4, 5, and 6, where the substrates coated with compositions containing hydroxyl-adjacent di-substituted phenolic anti-oxidants are compared directly to the coated substrate of the present invention). The presence of 2,6-di-tert-butyl-4-methyl-phenol in the coating compositions used to prepare the coated aluminum substrates of the present invention is critical to the filiform corrosion resistance performance of the invention. The examples further demonstrate the use of 2,6-di-tert-butyl-4-methyl-phenol within the claimed ranges of 0.5 to 10 percent by weight based on the total weight of resin solids in the powder coating composition.

(6) <u>ISSUES</u>

(i) No issues relating to 35 U.S.C. §112, first paragraph.

- (ii) No issues relating to 35 U.S.C. §112, second paragraph. The rejection of claim 4 under 35 U.S.C. §112, second paragraph, is now moot with the cancellation of claim 4.
- (iii) No issues relating to 35 U.S.C. §102. The rejections under 35 U.S.C. §102 are believed to be most with the incorporation of the subject matter of now cancelled claim 4 into independent claim 18.
- (iv) There are two issues relating to 35 U.S.C. §103(a).
- (a) Whether the subject matter of 5-7, 9, 13-15, and 18 are obvious in view of the teachings of Geary as evidenced by Laver taken in view of the teachings of Nakae.
- (b) Whether the subject matter of 5, 6, 13-15, and 18 are obvious in view of the teachings of Chasser as evidenced by Laver taken in view of the teachings of Nakae.

(7) GROUPING OF CLAIMS

Claims 5, 6, 13, 14 and 18 stand or fall together. Claims 7 and 9 stand or fall together. Claims 7 and 9 do not have Chasser applied against them (Issue (iv) (b)) and therefore stand separately from 5, 6, 13, 14 and 18. Claim 15 stands or falls alone. Claim 15 includes limitations that further distinguish this claim from the teachings of Geary, which supports the standing of claim 15.

- (8) ARGUMENTS
- (i) No issue.
- (ii) No issue.
- (iii) No issue.

(iv) (a) Whether the subj ct matter of 5-7, 9, 13-15, and 18 are obvious in view of the teachings of Geary as evidenced by Laver taken in view of the teachings of Nakae.

Claims 5, 6, 13, 14 and 18:

Claims 5, 6, 13, 14, and 18 are assumed to be rejected under 35 U.S.C. §103(a) as being obvious over Geary as evidenced by Laver and in view of Nakae. In both the Office Action of April 3, 2003, and the Final Office Action of August 26, 2003, the Examiner asserts that "Geary discloses thermosetting powder coating compositions comprising carboxylic acid-containing polyesters and beta-hydroxyalkylamide curing agents" (Geary, col. 1, lines 34-39). The Examiner further asserts that "examples show bis-hydroxyethylamide compounds and carboxylic acid group-containing polyester polymers having a number average molecular weight of 3500-4500. Example 1A suggests the combination of 72.4 percent by weight carboxylic acid-containing polyester, 4.3 percent by weight bis-hydroxyethylamide, and 1.5 percent by weight Irganox® 1076." Laver provides the structure of Irganox® 1076, which is n-octadecyl-3-(3,5-di-t-butyl-4-hydroxyphenyl)propionate, a phenolic compound having branched butyl groups in positions ortho to the hydroxy group. The coatings are applied to metal substrates, including aluminum. As noted by the Examiner in the Office Action of April 3, 2003, neither Geary nor Laver mentions the use of 2,6-di-tert-butyl-4-methyl-phenol. The Nakae reference teaches acid-epoxy type powder coating compositions comprising antioxidants having a melting point of from 50°C to 140°C. The Examiner relies on Nakae for teaching the use of 2,6-di-tert-butyl-4-methyl-phenol in powder coating compositions, asserting that certain phenol antioxidants, including 2,6-di-tert-butyl-4-methylphenol, are preferred in powder compositions because of their melting points. The Examiner concludes that it would have been obvious to one skilled in the art to choose 2,6-di-tert-butyl-4-methyl-phenol as a phenolic antioxidant in Geary's composition to form coatings having improved blocking resistance and melt processability.

Appellants respectfully submit that the basis on which the claims were rejected is not a valid rejection under 35 U. S. C. §103(a). First, if the references are considered each in their entirety, there is no motivation to combine the teachings of Geary with those of Nakae. Secondly, even if one were to combine the whole teaching of Geary with the whole teaching of Nakae, one would not arrive at the present invention as claimed.

References Considered as a Whole Offer No Motivation to Combine

The motivation to modify the prior art must flow from some teaching in the art that suggests the desirability or incentive to make the modification needed to arrive at the claimed invention. See In re Napier, 55 F.3d 610, 613, 34 U.S.P.Q.2d 1782, 1784 (Fed. Cir. 1995). Geary discloses thermosetting powder coating compositions that comprise a co-reactable particulate mixture of carboxylic acid group-containing polyesters and beta-hydroxyalkylamide curing agents (Abstract). Examples 1A, 2A, 3 and 4 disclose the use of Irganox® 1076 as an additive in these powder coating compositions. The additive is defined in footnote 3 of Example 1 as a polyphenol stabilizer that inhibits oxidation. Anti-oxidants are not taught, suggested, or recommended anywhere else in the reference. All of the examples in Geary demonstrate coatings applied over steel panels. Nakae discloses a different powder coating composition relying on a different cure chemistry, comprising an epoxy groupcontaining acrylic resin; a polycarboxylic acid; and an anti-oxidant having a melting point of from 50° to 140°C (Abstract). A list of over fifty anti-oxidants is provided (Nakae col. 3. Line 35-col. 4, line 49), which includes 2,6-di-tert-butyl-4-methyl-phenol and n-octadecyl-3-(3,5-di-t-butyl-4-hydroxyphenyl)propionate (Irganox® 1076). There is no teaching or suggestion in Nakae that the antioxidants used therein are suitable for use in any other coatings; moreover, coatings comprising carboxylic acid group-containing resins and betahydroxyalkylamide curing agents are not disclosed at all. The only coating composition disclosed in Nakae undergoes an acid-epoxy type cure mechanism. There is no teaching or suggestion that any of the anti-oxidants in Nakae would be effective in a carboxylic acid/beta-hydroxyalkylamide system such as that disclosed in Geary, much less that 2,6-di-tert-butyl-4-methyl-phenol would be equally effective as Irganox® 1076 in the system disclosed by Geary, such that one skilled in the art would be led to substitute the one anti-oxidant for the other.

The Examiner in both Office Actions alleges that the motivation to modify the composition of Geary with the 2,6-di-tert-butyl-4-methyl-phenol of Nakae is in order to form coatings having improved blocking resistance and melt processability. However, this proposed motivation to modify the Geary reference with the teaching of Nakae to use 2,6-di-tert-butyl-4-methyl-phenol is not supported. There is no teaching or suggestion in Nakae that 2,6-di-tert-butyl-4-methyl-phenol may be better than Irganox® 1076 for blocking resistance and melt processability in a coating composition that contains carboxylic acid functional polymers and beta-hydroxyalkylamide curing agents, such as that disclosed in Geary, or even that 2,6-di-tert-butyl-4-methyl-phenol is better than Irganox® 1076 for blocking resistance and melt processability in any powder coating composition.

There is absolutely no motivation to modify the compositions of Geary by particularly choosing the 2,6-di-tert-butyl-4-methyl-phenol of Nakae for improving blocking resistance and/or melt processability as suggested by the Examiner, or for any other reason, because none is offered by either reference, taken alone or in combination. Additionally, there is no teaching or suggestion in either reference, taken alone or in combination, to use the 2,6-di-tert-butyl-4-methyl-phenol of Nakae in a coating composition that contains carboxylic acid functional polymers and beta-hydroxyalkylamide curing agents, much less over an aluminum substrate, since neither reference demonstrates the use of such a combination over aluminum.

References Considered as a Whole Teach Away from Invention

If one were to combine the whole teachings of Geary with the whole teaching of Nakae, one would not arrive at the present invention as claimed; i. e., a coated aluminum substrate containing a cured coating derived from a coating composition comprising:

- (a) a polymer containing carboxylic functional groups;
- (b) a beta hydroxyalkylamide curing agent having functional groups reactive with the functional groups of the polymer which is present in an amount sufficient to cure the polymer; and
- (c) 0.5 to 10 weight percent based on the total weight of resin solids in the coating composition of 2,6-di-tert-butyl-4-methyl-phenol, the coated substrate being characterized as having improved filiform corrosion resistance compared to a similar coated substrate that does not contain (c) in the cured coating.

"It is impermissible within the framework of section 103 to pick and choose from any one reference only so much of it as will support a given position, to the exclusion of other parts necessary to the full appreciation of what such reference fairly suggests to one of ordinary skill in the art." In re Wesslau, 353 F.2d 238, 241, 147 U.S.P.Q. 391, 393 (C.C.P.A. 1965). See also MPEP §2141.02: "[a] prior art reference must be considered in its entirety, i. e., as a whole, including portions that would lead away from the claimed invention." If the references are considered each in their entirety, the teachings of Nakae combined with Geary teach away from the present invention. As mentioned earlier, Geary discloses thermosetting powder coating compositions that comprise a co-reactable particulate mixture of carboxylic acid groupcontaining polyesters and beta-hydroxyalkylamide curing agents. discloses a powder coating composition comprising an epoxy group-containing acrylic resin; a polycarboxylic acid; and any of over fifty anti-oxidants having a melting point of from 50° to 140°C. While a variety of metal substrates including both steel and aluminum are disclosed in Nakae (col. 6, lines 26-30), the only substrate on which the coating composition is demonstrated is steel. Likewise, all of the examples in Geary demonstrate coatings applied over steel panels. If one were to combine both references after considering them in their entirety, in order to prepare a coated substrate containing a cured coating, which in turn is derived from a coating composition comprising a polymer containing carboxylic functional groups, a beta-hydroxyalkylamide curing agent, and an anti-oxidant, one skilled in the art would use a steel substrate, since both references clearly demonstrate the use of their respective coating compositions over steel. Additionally, one is not likely to choose 2,6-di-tert-butyl-4-methyl-phenol from among a list of over fifty other anti-oxidants, without any guidance to do so offered by the Nakae reference. Further, since neither reference recognizes the improvement in filiform corrosion resistance over aluminum afforded by the use of 2,6-di-tert-butyl-4-methyl-phenol, there would be no particular impetus to use an aluminum substrate or 2,6-di-tert-butyl-4-methyl-phenol, and therefore it would not have been obvious to do so.

Claim 15

The discussion under the heading "Claims 5, 6, 13, 14, and 18" is incorporated herein by reference.

Claim 15 recites the coated aluminum substrate of claim 18 where said polymer is an acrylic polymer. The Examiner asserts in both Office Actions that Example 2A in Geary "shows the combination of carboxylic acid-containing polyester, carboxylic acid-containing acrylic polymer, a bis-hydroxyethylamide, and Irganox® 1076". As noted above, Geary discloses thermosetting powder coating compositions that comprise a co-reactable particulate mixture of carboxylic acid group-containing *polyesters* and beta-hydroxyalkylamide curing agents. The acrylic polymer disclosed in Example 2 is used in very minor amounts (less than 5 percent by weight of the resinous binder; less than 3.5 percent by weight of total resin solids). One skilled in the art would hardly look to Geary for a teaching of coating compositions containing carboxylic acid functional acrylics because Geary teaches a powder coating composition containing polyester polymers. Claim 15 is therefore believed to further distinguish the coated aluminum substrate of the present invention from Geary taken alone or in combination with Nakae.

Claims 7 and 9

The discussion under the heading "Claims 5, 6, 13, 14, and 18" is incorporated herein by reference.

Claim 7 recites the coated aluminum substrate of claim 18 wherein said polymer has an equivalent weight of from 200 to 2500. Claim 9 recites the

coated aluminum substrate of claim 18 wherein the beta-hydroxyalkylamide is bis-hydroxyethylamide. The arguments presented above with respect to claims 5, 6, 13, 14, and 18 apply here, and it is noted as above in the Grouping of Claims that claims 7 and 9 stand or fall together, independently of 5, 6, 13, 14 and 18. Claims 7 and 9 do not have Chasser applied against them (Issue (iv)(b)) and therefore stand separately from 5, 6, 13, 14 and 18.

(iv)(b) Whether the subject matter of 5, 6, 13-15, and 18 are obvious in view of the teachings of Chasser as evidenced by Laver taken in view of the teachings of Nakae.

Claims 5, 6, 13, 14 and 18:

Claims 5, 6, 13, 14, and 18 are assumed to be rejected under 35 U.S.C. §103(a) as being obvious over Chasser as evidenced by Laver and in view of Nakae. In both the Office Action of April 3, 2003, and the Final Office Action of August 26, 2003, the Examiner asserts that "Chasser discloses curable powder coating compositions comprising polymers with functional groups and curing agents reactive with the polymers that are applied to aluminum substrates (Abstract)." The Examiner further asserts that the polymers are selected from carboxylic acid group-containing polyester polymers having a number average molecular weight of 2000-3000, carboxylic acid group-containing acrylic polymers, carboxylic acid group-containing polyurethane polymers, and epoxy functional polymers. Polymers are used in preferred amounts of 50-85 percent beta-hydroxyalkylamides Curing agents include by weight. triglycidylisocyanurate (TGIC). Irganox® anti-oxidants are noted as suitable additives. Example A shows the combination of a carboxylic acid-containing polyester, 12.9 percent by weight TGIC, and about 1 percent by weight Irganox® 1076. Laver provides the structure of Irganox® 1076. Example C shows the combination of a carboxylic acid-containing polyester, a betahydroxyalkylamide curing agent, and Tinuvin® 900, a phenolic ultraviolet light stabilizer having substituents at ortho positions to the hydroxy group. As noted by the Examiner in the Office Action of April 3, 2003, neither Chasser nor Laver mentions the use of 2,6-di-tert-butyl-4-methyl-phenol. The Nakae reference teaches acid-epoxy type powder coating compositions comprising antioxidants having a melting point of from 50°C to 140°C. The Examiner relies on Nakae for teaching the use of 2,6-di-tert-butyl-4-methyl-phenol in powder coating compositions, asserting that certain phenol antioxidants, including 2,6-di-tert-butyl-4-methyl-phenol, are preferred in powder compositions because of their melting points. The Examiner concludes that it would have been obvious to one skilled in the art to choose 2,6-di-tert-butyl-4-methyl-phenol as a phenolic antioxidant in Chasser's composition to form coatings having improved blocking resistance and melt processability.

Appellants respectfully submit that the basis on which the claims were rejected is not a valid rejection under 35 U. S. C. §103(a). First, if the references are considered each in their entirety, there is no motivation to combine the teachings of Chasser with those of Nakae. Secondly, even if one were to combine the whole teaching of Chasser with the whole teaching of Nakae, one would not arrive at the present invention as claimed.

References Considered as a Whole Offer No Motivation to Combine

The Examiner in both Office Actions alleges that the motivation to modify the composition of Chasser with the 2,6-di-tert-butyl-4-methyl-phenol of Nakae is in order to form coatings having improved blocking resistance and melt processability. However, this proposed motivation to modify the Chasser reference with the teaching of Nakae to use 2,6-di-tert-butyl-4-methyl-phenol is not supported. As noted above, there is *no* teaching or suggestion in Nakae that 2,6-di-tert-butyl-4-methyl-phenol is better than Irganox® 1076 for blocking resistance and melt processability in *any* powder coating composition, let alone that 2,6-di-tert-butyl-4-methyl-phenol may be better than Irganox® 1076 for blocking resistance and melt processability in a coating composition that contains carboxylic acid functional polymers and beta-hydroxyalkylamide curing agents, which is only one of a collection of coating systems disclosed by Chasser that includes carboxylic acid functional polymers/polyepoxide curing

agents, carboxylic acid functional polymers/TGIC, and epoxy functional polymers/polyacid curing agents.

One skilled in the art would not be motivated to improve blocking resistance and/or melt processability of the Chasser composition by particularly choosing the 2,6-di-tert-butyl-4-methyl-phenol of Nakae because no teaching or suggestion is made in either reference, taken alone or in combination, of improved blocking resistance and/or melt processability offered by 2,6-di-tert-butyl-4-methyl-phenol. Additionally, there is no teaching or suggestion in either reference, taken alone or in combination, to select the 2,6-di-tert-butyl-4-methyl-phenol of Nakae for any reason and use it in a coating composition that contains carboxylic acid functional polymers and beta-hydroxyalkylamide curing agents.

References Considered as a Whole Teach Away from Invention

If one were to combine the whole teachings of Chasser with the whole teaching of Nakae, one would not arrive at the present invention as claimed. Chasser discloses curable powder coating compositions comprising polymers with functional groups and curing agents reactive with the polymers. The composition further comprises an oligomeric acid functional polyester with a particular structure as shown in col. 2, lines 40-48, which is the defining feature The oligomeric polyester provides the of the Chasser composition. composition with improved adhesion to metal and filiform corrosion resistance. The polymers with functional groups used by Chasser may be any of a variety of polymers with any of a variety of functional groups. Likewise, the curing agents may be selected from any of a number of different materials, depending on the functionality of the polymer. As mentioned above, carboxylic acid functional polymers/polyepoxide curing agents, carboxylic acid functional polymers/TGIC, and epoxy functional polymers/polyacid curing agents are all Nakae discloses disclosed by Chasser as suitable coating systems. exclusively an epoxy group-containing acrylic resin; a polycarboxylic acid; and an anti-oxidant having a melting point of from 50° to 140°C (Abstract). As noted above, list of over fifty anti-oxidants is provided, which includes 2,6-ditert-butyl-4-methyl-phenol and Irganox® 1076. There is no teaching or suggestion in Nakae that the anti-oxidants used therein are suitable for use in any other coatings; moreover, coatings comprising carboxylic acid group-containing resins and beta-hydroxyalkylamide curing agents are not disclosed at all in Nakae. The only coating composition disclosed in Nakae undergoes an acid-epoxy type cure mechanism. There is no teaching or suggestion that any of the other anti-oxidants in Nakae would be effective in a carboxylic acid/beta-hydroxyalkylamide system, much less that 2,6-di-tert-butyl-4-methyl-phenol would be equally effective as Irganox® 1076 a carboxylic acid/beta-hydroxyalkylamide system, such that one skilled in the art would be led to use the particular anti-oxidant 2,6-di-tert-butyl-4-methyl-phenol in the particular coating composition comprising carboxylic acid group-containing resins and beta-hydroxyalkylamide curing agents.

If the references are considered each in their entirety, the teachings of Nakae combined with Chasser teach away from the present invention. As mentioned earlier, Chasser discloses curable powder coating compositions comprising polymers with any of a number of functional groups and curing Nakae discloses a powder coating agents reactive with the polymers. an epoxy group-containing acrylic resin; a composition comprising polycarboxylic acid; and any of over fifty anti-oxidants having a melting point of from 50° to 140°C. If one were to combine both references after considering them in their entirety, in order to prepare a coated substrate containing a cured coating, which in turn is derived from a coating composition comprising a polymer containing functional groups, a curing agent, and an anti-oxidant, one skilled in the art would use an epoxy group-containing resin and a polycarboxylic acid curing agent, since that is the coating composition common to both references. Additionally, one is not likely to choose 2,6-di-tert-butyl-4methyl-phenol from among a list of over fifty other anti-oxidants, without any guidance to do so offered by the Nakae reference. Further, since neither reference recognizes the improvement in filiform corrosion resistance over aluminum afforded by the use of 2,6-di-tert-butyl-4-methyl-phenol (Chasser has already achieved filiform corrosion resistance with the oligomeric polyester),

there would be no particular reason to use 2,6-di-tert-butyl-4-methyl-phenol, and therefore it would not have been obvious to do so.

Claim 15

The discussion under the heading "Claims 5, 6, 13, 14, and 18" is incorporated herein by reference.

Claim 15 recites the coated aluminum substrate of claim 18 where said polymer is an acrylic polymer. The arguments presented above with respect to claims 5, 6, 13, 14, and 18 apply here, and it is noted as above in the Grouping of Claims that claim 15 includes limitations that further distinguish this claim from the teachings of Geary (Issue (iv)(a)), which supports the standing of claim 15.



CONCLUSION

Appellants have shown that if the references are considered each in their entirety, there is no motivation to combine the teachings of the Geary reference with Nakae in order to arrive at the present invention. Likewise, there is no motivation to combine the teachings of the Chasser reference with Nakae in order to arrive at the present invention. Appellants have further shown that if one were to combine the whole teachings of Geary or Chasser with the whole teaching of the Nakae, one would not arrive at the present invention as claimed.

Based on the above, Appellants respectfully request that the Board reverse the Examiner on the rejections of claims 5, 6, 7, 9, 13, 14, 15 and 18 as unpatentable under 35 U.S.C. 103 over the applied prior art and allow these claims.

Respectfully submitted,

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<u>Appendix</u>

THE CLAIMS ON APPEAL

- 5. The coated aluminum substrate of claim 18 wherein said polymer containing carboxylic groups is selected from the group consisting of acrylic polymers, polyester polymers, and polyurethane polymers.
- 6. The coated aluminum substrate of claim 18 wherein said polymer has a number average molecular weight of from 1,000 to 20,000.
- 7. The coated aluminum substrate of claim 18 wherein said polymer has an equivalent weight equal from 200 to 2,500.
- 9. The coated aluminum substrate of claim 18 wherein the betahydroxyalkylamide is bis-hydroxyethylamide.
- 13. The coated aluminum substrate of claim 18 wherein said polymer is present in an amount ranging from 10 to 80 weight percent based on the total weight resin solids in the powder coating composition.
- 14. The coated aluminum substrate of claim 18 wherein said curing agent is present in an amount ranging from 2 to 40 weight percent based on the total weight resin solids in the powder coating composition.

- 15. The coated aluminum substrate of claim 18 where said polymer is an acrylic polymer.
- 18. A coated aluminum substrate containing a cured coating derived from a coating composition comprising:
 - a polymer containing carboxylic functional groups;
- b. a beta hydroxyalkylamide curing agent having functional groups reactive with the functional groups of the polymer which is present in an amount sufficient to cure the polymer; and
- c. 0.5 to 10 weight percent based on the total weight of resin solids in the coating composition of 2,6-di-tert-butyl-4-methyl-phenol, the coated substrate being characterized as having improved filiform corrosion resistance compared to a similar coated substrate that does not contain (c) in the cured coating.